Present Claims 1-3 and 5-19 relate to an isolated coryneform bacterium in which L-arginine biosynthesis is not repressed by an *argR* gene in a normal manner, and which has L-arginine producing ability. The inventors have surprisingly found that the presently claimed bacteria are especially useful for producing L-arginine.

The rejection of Claims 1-3 under 35 U.S.C. § 101 as being directed toward non-statutory subject matter has been obviated by appropriate amendment. As the Examiner will note, Claim 1 has been amended to replace "A coryneform bacterium" with "An isolated coryneform bacterium." Accordingly, the rejection is no longer tenable and should be withdrawn.

The rejection of Claims 1-3 under 35 U.S.C. § 112, first paragraph, is respectfully traversed. In the Official Action, the position is taken that Claims 1-3 contain subject matter which was not described in the specification in such a way as to reasonably convey to one skilled in the relevant art that the inventor, at the time the application was filed, had possession of the claimed invention. However, Applicants would like to direct the Examiner's attention to the following points.

Some *argR* genes of coryneform bacteria were well known and available at the time the application was filed. For example, the *argR* gene of *Corynebacterium glutamicum* was registered in Genbank as accession No. AF041436. The nucleotide sequence of an *argR* gene for *Mycobacterium tuberculosis* was disclosed in Nature, vol. 393, p. 537 (1998). In this regard, Table III-4-1 of Identification Manual of Actinomycetes (published by Business Center for Academic Societies Japan) (copy attached as Exhibit A) shows that *Mycobacterium* belongs to the same suborder as *Corynebacterium*.

It had not been known whether a protein encoded by argR gene actually functions as a repressor of L-arginine biosynthetic system in coryneform bacteria. However, the inventors of the present invention have now discovered, for the first time, an argR gene in Brevibacterium flavum, which belongs to coryneform bacteria. Therefore, other coryneform bacteria are also reasonably now expected to have argR genes, and proteins encoded by argR genes are reasonably expected to function as a repressor of the L-arginine biosynthetic system. Consequently, Applicants submit that the specification also provides sufficient enablement for the claimed invention even in such cases when using argR gene other than that of B. flavum.

For these reasons, the rejection should be withdrawn.

The rejection of Claims 1-3 under 35 U.S.C. §112, second paragraph, has been obviated by appropriate amendment. In particular, Claims 1 and 2 have been amended by replacing "arginine repressor does not function in a normal manner" with "L-arginine biosynthesis is not repressed by an *argR* gene in a normal manner."

Claim 2 has also been amended to replace "a gene coding for the arginine repressor" with "the *argR* gene which has the nucleotide sequence shown in SEQ ID NO:17 or has such a degree of homology that it should cause homologous recombination with the nucleotide sequence shown in SEQ ID NO:17." As stated above, it is now reasonably expected that other coryneform bacteria than *B. flavum* have *argR* genes. The *argR* genes are expected to have the sequences homologous to the sequence of SEQ ID NO:17. Therefore, Applicants believe that these *argR* genes can be disrupted by homologous recombination and this rejection should be overcome.

Claim 3 has been amended by replacing "an amino acid sequence showing homology to the amino acid sequence" with "an amino acid sequence which is encoded by an argR gene having such a degree of homology that it should cause homologous recombination with the argR gene coding to the amino acid sequence shown in SEQ ID NO:18."

Accordingly, the rejection is no longer tenable and should be withdrawn.

Applicants expressly state on the record that none of these amendments were made or necessary to distinguish the present claims from the prior art.

Applicants submit that the application is now in condition for allowance, and early notification of such action is earnestly solicited.

Respectfully submitted,

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Amendment Filed on:

12.28.01

IN THE CLAIMS

Please amend the claims as follows:

- --1. (Amended) [A] <u>An isolated</u> coryneform bacterium in which [an arginine repressor does not function] <u>L-arginine biosynthesis is not repressed by an *argR* gene in a normal manner, and which has L-arginine producing ability.</u>
- 2. (Amended) The <u>isolated</u> coryneform bacterium according to Claim 1, wherein [the arginine repressor does not function in a normal manner due to disruption of a gene coding for the arginine repressor] <u>L-arginine biosynthesis is not repressed by the *argR* gene in a normal manner due to disruption of the *argR* gene which has the nucleotide sequence shown in SEQ ID NO:17 or has such a degree of homology that it should cause homologous recombination with the nucleotide sequence shown in SEQ ID NO:17, and which is on a chromosome of the bacterium.</u>
- 3. (Amended) The <u>isolated</u> coryneform bacterium according to Claim 2, wherein the [arginine repressor has] <u>argR</u> gene encodes the amino acid sequence shown in SEQ ID NO:18 or an amino acid sequence [showing homology to the amino acid sequence] <u>which is encoded by an argR gene having such a degree of homology that it should cause homologous</u>

recombination with the argR gene coding for the amino acid sequence shown in SEQ ID

NO:18.--

Please add the following new claims:

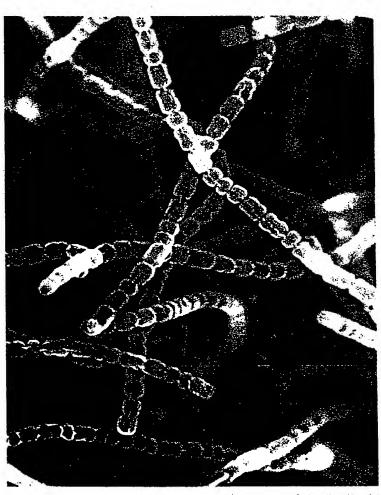
--5. (New) to 19. (New)--

EXHIBIT

放線菌の 類と同分

日本放線菌学会 編

edited by The Society for Actinomycetes Japan



Streptomyces griseus (ストレプトマイシン生)

財団法人 日本学会事務センター



M. 7032.

定価はカバーに表示

放線菌の分類と同定 Identification Manual of Actinomycetes

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〈無断複写・転載を禁ず〉 Printed in Japan

Family Corynebacteriaceae and related families

= 鈴木健一朗 :

本菌群は病原菌として古くから知られている桔核菌 (Mycobacterium buberculosis) やジフテリア菌 (Corynebacterium diphtheriae) を含み、Nocardia 風と合わせて各 属の頭文字を取り、CMN グループと呼ばれ、とくに医学 細菌学上重要な菌群とされてきた。その後,グルタミン酸 生産图 Corynebacteirum glutamicum がこの仲間に入り、 アミノ酸発酵の主役となると、産業微生物学的にも注目さ れるようになり, さらに Rhodococcus 属には多くの難分解 性人工有機化合物の分解能力が知られるようになって環境 微生物学上の重要性も増大した。Nocardia 属は生長に伴 って断裂する菌糸を持ち、かつては多くの放線菌を形態だ けで Nocardia と称し、その不均一性が問題となっていた が,1970年代の化学分類学的研究によって整理され,ほぼ 現在の Nocardia 属の概念が足若した。そしてミコール酸 含有という共通の性質を持つ函群は 16S rDNA の解析に よっても支持され,現在は表 III-4-1 に示す 10 属を数え る。これらの各属は 16SrDNA による系統では Nocardia と Rhodococcus が近縁で同じ科に入るほかは、それぞれほぼ 科レベルで分岐している (Rainey et al., 1995, Stackebrandt et al., 1997)。ミコール酸は2-アルキル3-ヒドロキシ脂肪 酸という―般構造を持つ,分枝ヒドロキシ脂肪酸である。 とくに Mycobacterium 属のミコール酸はその秘炭素数が On it is about a set may respect to

90 にも達する特異な脂肪酸で、その炭素数の分布幅は分類詳に特徴的である。二度結合数と官能基の有無にも分類詳ごとの特徴が見られる(第 II 部第 3 章第 7 節ミコール酸参照)。本章の対象は、Stackebrandt et al. (1997) の高次数 III-4-1. Suborder Corynebacterineae	
Corynebacteriaceae	Corynebacterium
	Turicella
Dietziaceae	Dietzia
Gordoniaceae	Gordonia
Mycobacteriaceae	Mycobacterium
Nocardiaceae	Nocardia
	Rhodococcus
Tsukamurellaceae	Tsukamurella
帰属不明	Skermania

Williamsia

分類群では、order Actinomycetales の suborder Corynebacterineae に対応する。本菌群の細菌の細胞壁は mesodiaminopimelic acid (A₂pm) 直結型のペプチドグリカン (Aly) と、その外側に arabinose と galactose からなる多 糖層を持つ点が共通である。そしてミコール酸はアラビノ ガラクタンに共有結合し、あるいはトレハロース 6,6'ジ ミコール酸のような糖脂質として存在する。この函群の中 にも例外的にミコール酸を持たない分類群も存在する。 Turicella 民と Corynebacterium amycolatum である。他の 放線菌群と同様、化学分類学的表現性状と系統分類はよく 対応しているが、表現性状だけで属を特定するのが困難な 場合もあるので、分類および同定には統合的な判断が求め られる。この菌群においてとくに有効な属の識別指標はメ ナキノンの分子種 (Collins et al., 1977) と、グリコリル テストによる細胞壁アシル型,そしてリン脂質とミコール 酸である。これらの化学分類学的性状による属の概念を表 Ⅲ-4-2 に示す。〕

本函群の生理・生化学性状は、とくに炭素化合物の資化性が多様で、種の特徴の記載には重要である。臨床株が主体の Corynebacterium, Mycobacterium, Turicella を除き、一般には Gordon の方法 (Gordon & Mihm, 1957)、または Goodfellow の方法 (Goodfellow, 1971) が用いられる。

本菌群は応用分野から臨床分野まで幅広い研究対象となり、人間とのかかわりも多様である。したがって、目的に応じた微生物との接し万があってしかるべきであり、微生物の認識の仕方も様々である。遺伝子解析技術を便った分類体系の構築に対して、同じデータベースを用いた検出手法とは相互に評価して一層価値が高まるものである。医学微生物には重要な目的が含まれている。一般微生物の分類を専門とする立場で本章を執筆しているので、病原放線菌の同定を行うときには医学微生物学の専門寄も合わせて参考にして、重要な目的を見失わないようにされることを願うものである。

Corynebacterium **医**

本属は基準種 C. diphtheriae に代表される多くの人奇病

